

# Labiprofen 150 mg/ml solution for injection

Authorised

- Ketoprofen

## Product identification

**Medicine name:**

Labiprofen 150 mg/ml solution for injection

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**Active substance:**

Ketoprofen

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**Target species:**

Pig  
Cattle  
Horse

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**Route of administration:**

Intramuscular use  
Intravenous use

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## Product details

**Active substance and strength:**

Ketoprofen  
150.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Pig**

- Meat and offal. 3 day

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**Cattle**

- Meat and offal. 2 day

- Milk. 0 hour

**Intravenous use:**

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**Horse**

- Meat and offal. 1 day

- Milk. no withdrawal period

Milk: Not authorised for use in mares producing milk for human consumption.

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**Cattle**

- Meat and offal. 2 day

- Milk. 0 hour

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM01AE03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Sweden

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**Available in:**

Sweden

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**Package description:**

box containing 10 vials of 250 ml

box containing 10 vials of 100 ml

box containing 12 vials of 50 ml

box containing 1 vial of 250 ml

box containing 1 vial of 100 ml

box containing 1 vial of 50 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Labiana Life Sciences S.A.

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**Marketing authorisation date:**

4/03/2024

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**Manufacturing sites for batch release:**

Labiana Life Sciences S.A.

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**Responsible authority:**

Swedish Medical Products Agency

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**Authorisation number:**

64609

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**Date of authorisation status change:**

4/03/2024

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0388/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland  
France Germany Greece Hungary Ireland Italy Latvia Lithuania  
Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Sweden  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents